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OM protein - protein search, using sw model

Run on: August 31, 2004, 19:53:21 ; Search time 32 Seconds
(without alignments)
296.849 Million cell updates/sec

Title: US-09-589-777C-2
Perfect score: 968
Sequence: 1 HTHQDFQPVHLVALNTPLS.....CHNSYIVLCIENSFMTSFSK 184

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 389414 seqs, 51625971 residues

Total number of hits satisfying chosen parameters: 389414

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA: *
1: /cgn2_6/ptodata/2/iaa/5A_COMB.pep.*
2: /cgn2_6/ptodata/2/iaa/5B_COMB.pep.*
3: /cgn2_6/ptodata/2/iaa/6A_COMB.pep.*
4: /cgn2_6/ptodata/2/iaa/6B_COMB.pep.*
5: /cgn2_6/ptodata/2/iaa/PCTUS_COMB.pep.*
6: /cgn2_6/ptodata/2/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	968	100.0	191	4	US-09-561-500-13
2	968	100.0	191	4	US-09-561-108-13
3	968	100.0	191	4	US-09-561-526-13
4	968	100.0	191	4	US-09-561-499-13
5	968	100.0	191	4	US-09-998-831-13
6	965	99.7	195	1	US-08-159-784-2
7	916	94.6	185	3	US-08-985-526-36
8	840	86.8	182	4	US-09-561-500-14
9	840	86.8	182	4	US-09-561-108-14
10	840	86.8	182	4	US-09-315-689-3
11	840	86.8	182	4	US-09-561-526-14
12	840	86.8	182	4	US-09-561-499-14
13	840	86.8	182	4	US-09-998-831-14
14	840	86.8	183	3	US-09-206-059-2
15	822	84.9	178	4	US-09-315-689-5
16	550	56.8	191	1	US-08-159-784-3
17	273.5	28.3	124	4	US-09-231-077D-10
18	198	20.5	123	4	US-09-231-077D-11
19	150	15.5	35	3	US-09-046-985-2
20	150	15.5	35	3	US-09-474-743-2
21	108	11.2	20	2	US-08-740-168A-1
22	108	11.2	20	3	US-09-349-429-1
23	108	11.2	20	4	US-09-315-689-1
24	108	11.2	20	4	US-09-174-282-1
25	108	11.2	20	4	US-09-154-302-1
26	101	10.4	16	3	US-09-385-442-32
27	101	10.4	22	3	US-09-046-985-7
					Sequence 13, Appl
					Sequence 13, Appl
					Sequence 13, Appl
					Sequence 13, Appl
					Sequence 13, Appl
					Sequence 2, Appl
					Sequence 36, Appl
					Sequence 14, Appl
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					Sequence 3, Appl
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					Sequence 14, Appl
					Sequence 2, Appl
					Sequence 5, Appl
					Sequence 3, Appl
					Sequence 10, Appl
					Sequence 11, Appl
					Sequence 2, Appl
					Sequence 2, Appl
					Sequence 1, Appl
					Sequence 1, Appl
					Sequence 1, Appl
					Sequence 1, Appl
					Sequence 32, Appl
					Sequence 7, Appl

28	101	10.4	22	3	US-09-474-743-7	Sequence 7, Appl
29	82	8.5	15	4	US-09-231-077D-12	Sequence 12, Appl
30	79	8.2	682	4	US-09-489-039A-10595	Sequence 10595, A
31	79	8.2	1112	4	US-09-717-364A-27	Sequence 27, Appl
32	77	8.0	439	4	US-09-252-991A-19623	Sequence 19623, A
33	76.5	7.9	1050	4	US-09-428-711A-16	Sequence 16, Appl
34	76	7.9	506	4	US-09-252-991A-18165	Sequence 18165, A
35	75	7.7	190	3	US-09-046-985-15	Sequence 15, Appl
36	75	7.7	190	3	US-09-474-743-15	Sequence 15, Appl
37	75	7.7	587	2	US-08-871-266B-18	Sequence 18, Appl
38	75	7.7	587	2	US-09-018-864A-18	Sequence 18, Appl
39	75	7.7	587	3	US-08-871-267B-24	Sequence 24, Appl
40	75	7.7	587	3	US-09-618-419-24	Sequence 24, Appl
41	74.5	7.7	577	2	US-08-756-317-13	Sequence 13, Appl
42	74	7.6	15	4	US-09-231-077D-13	Sequence 13, Appl
43	74	7.6	1646	4	US-09-252-991A-22312	Sequence 22312, A
44	74	7.6	6396	4	US-09-410-551B-72	Sequence 72, Appl
45	73.5	7.6	443	4	US-09-252-991A-26460	Sequence 26460, A

ALIGNMENTS

RESULT 1
US-09-561-500-13
; Sequence 13, Application US/09561500
; Patent No. 6342219
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY COMPOSITIONS FOR SELECTIVELY INHIBITING VEGF
; FILE REFERENCE: 4001.002500
; CURRENT APPLICATION NUMBER: US/09/561,500
; CURRENT FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/131,432
; PRIOR FILING DATE: 1999-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
; LENGTH: 191
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-561-500-13

Query Match 100.0%; Score 968; DB 4; Length 191;
Best Local Similarity 100.0%; Pred. No. 3.8e-112;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	HTHODFQPVHLVALNTPLSGMGRGIRGADFCQFQQAARAVGLSGTFRFLSSRLQDLYSI	60
Db	8	HTHODFQPVHLVALNTPLSGMGRGIRGADFCQFQQAARAVGLSGTFRFLSSRLQDLYSI	67
QY	61	VRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSGDGRDVLRHPAPQKSVW	120
Db	68	VRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSGDGRDVLRHPAPQKSVW	127
QY	121	HGSDPSGRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT	180
Db	128	HGSDPSGRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT	187
QY	181	SFSK 184	
Db	188	SFSK 191	

RESULT 2
US-09-561-108-13
; Sequence 13, Application US/09561108
; Patent No. 6342221
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe

APPLICANT: Rolf A. Brekken
TITLE OF INVENTION: ANTIBODY CONJUGATE COMPOSITIONS FOR SELECTIVELY INHIBITING VEGF
FILE REFERENCE: 4001.002584
CURRENT APPLICATION NUMBER: US/09/561,108
CURRENT FILING DATE: 2000-04-28
PRIOR APPLICATION NUMBER: 60/131,432
PRIOR FILING DATE: 1999-04-28
NUMBER OF SEQ ID NOS: 44
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 13
LENGTH: 191
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-561-108-13

Query Match 100.0%; Score 968; DB 4; Length 191;
Best Local Similarity 100.0%; Pred. NO. 3.8e-112;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 67
QY 61 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 120
Db 68 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 127
QY 121 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 128 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 187
QY 181 SFSK 184
Db 188 SFSK 191

RESULT 3
US-09-561-526-13
Sequence 13, Application US/09561526
Patent No. 6416758
GENERAL INFORMATION:
APPLICANT: Philip E. Thorpe
APPLICANT: Rolf A. Brekken
TITLE OF INVENTION: ANTIBODY CONJUGATE KITS FOR SELECTIVELY INHIBITING VEGF
FILE REFERENCE: 4001.002586
CURRENT APPLICATION NUMBER: US/09/561,526
CURRENT FILING DATE: 2000-04-28
PRIOR APPLICATION NUMBER: 60/131,432
PRIOR FILING DATE: 1999-04-28
NUMBER OF SEQ ID NOS: 44
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 13
LENGTH: 191
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-561-526-13

Query Match 100.0%; Score 968; DB 4; Length 191;
Best Local Similarity 100.0%; Pred. NO. 3.8e-112;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 67
QY 61 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 120
Db 68 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 127

QY 121 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 128 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 187
QY 181 SFSK 184
Db 188 SFSK 191

RESULT 4
US-09-561-499-13
Sequence 13, Application US/09561499
Patent No. 6524583
GENERAL INFORMATION:
APPLICANT: Philip E. Thorpe
APPLICANT: Rolf A. Brekken
TITLE OF INVENTION: ANTIBODY METHODS FOR SELECTIVELY INHIBITING VEGF
FILE REFERENCE: 4001.002582
CURRENT APPLICATION NUMBER: US/09/561,499
CURRENT FILING DATE: 2000-04-28
PRIOR APPLICATION NUMBER: 60/131,432
PRIOR FILING DATE: 1999-04-28
NUMBER OF SEQ ID NOS: 44
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 13
LENGTH: 191
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-561-499-13

Query Match 100.0%; Score 968; DB 4; Length 191;
Best Local Similarity 100.0%; Pred. NO. 3.8e-112;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 67
QY 61 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 120
Db 68 VRRADGSGVPIVNLKDEVLSFSGSQGLQPGARIFSDGRDVLRLHPAPQKSVW 127
QY 121 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 128 HGSDPSGRRRLMESYCETWRTTGTGATQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 187
QY 181 SFSK 184
Db 188 SFSK 191

RESULT 5
US-09-998-831-13
Sequence 13, Application US/09998831
Patent No. 6676941
GENERAL INFORMATION:
APPLICANT: Philip E. Thorpe
APPLICANT: Rolf A. Brekken
TITLE OF INVENTION: ANTIBODY CONJUGATE COMPOSITIONS FOR SELECTIVELY
TITLE OF INVENTION: INHIBITING VEGF
FILE REFERENCE: 4001.002584
CURRENT APPLICATION NUMBER: US/09/998,831
CURRENT FILING DATE: 2001-11-30
PRIOR APPLICATION NUMBER: 09/561,108
PRIOR FILING DATE: 2000-04-28
NUMBER OF SEQ ID NOS: 44
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 13
LENGTH: 191
TYPE: PRT
ORGANISM: Artificial Sequence

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; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-998-831-13

Query Match      100.0%; Score 968; DB 4; Length 191;
Best Local Similarity 100.0%; Pred. No. 3.8e-112;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRFLSSRLQDLYSI 67

QY 61 VRRADRGSPVIVNLKDEVLSLSPSWDSLFSGSQGQLQPGARIFSDGRDVLHRHPAWPKSVW 120
Db 68 VRRADRGSPVIVNLKDEVLSLSPSWDSLFSGSQGQLQPGARIFSDGRDVLHRHPAWPKSVW 127

QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 180
Db 128 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 187

QY 181 SFSK 184
Db 188 SFSK 191

RESULT 6
US-08-159-784-2
; Sequence 2, Application US/08159784
; Patent No. 5643783
; GENERAL INFORMATION:
; APPLICANT: Bjorn R. Olsen
; TITLE OF INVENTION: NOVEL COLLAGEN AND USES THEREOF
; NUMBER OF SEQUENCES: 9
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Fish & Richardson
; STREET: 225 Franklin Street
; CITY: Boston
; STATE: Massachusetts
; COUNTRY: U.S.A.
; ZIP: 02110-2804
; COMPUTER READABLE FORM:
; MEDIUM TYPE: 3.5" Diskette, 1.44 Mb
; COMPUTER: IBM PS/2 Model 50Z or 55SX
; OPERATING SYSTEM: MS-DOS (Version 5.0)
; SOFTWARE: WordPerfect (Version 5.1)
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/159,784
; FILING DATE: December 1, 1993
; CLASSIFICATION: 530
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: John F. Freeman
; REGISTRATION NUMBER: 29,066
; REFERENCE/DOCKET NUMBER: 00246/170001
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617) 542-5070
; TELEFAX: (617) 542-8906
; TELEX: 200154
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 195
; TYPE: amino acid
; STRANDEDNESS: N/A
; TOPOLOGY: N/A
US-08-159-784-2

Query Match      99.7%; Score 965; DB 1; Length 195;
Best Local Similarity 99.5%; Pred. No. 9.2e-112;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRFLSSRLQDLYSI 60
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Db 12 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRFLSSRLQDLYSI 71
QY 61 VRRADRGSPVIVNLKDEVLSLSPSWDSLFSGSQGQLQPGARIFSDGRDVLHRHPAWPKSVW 120
Db 72 VRRADRGSPVIVNLKDEVLSLSPSWDSLFSGSQGQVQPGARIFSDGRDVLHRHPAWPKSVW 131
QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 180
Db 132 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 191
QY 181 SFSK 184
Db 192 SFSK 195

RESULT 7
US-08-985-526-36
; Sequence 36, Application US/08985526
; Patent No. 6080728
; GENERAL INFORMATION:
; APPLICANT: Mixson, James A
; TITLE OF INVENTION: CARRIER:DNA COMPLEXES CONTAINING DNA
; TITLE OF INVENTION: ENCODING ANTI-ANGIOGENIC PEPTIDES AND THEIR USE IN GENE
; TITLE OF INVENTION: THERAPY
; NUMBER OF SEQUENCES: 43
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Connolly, Bove, Lodge, & Hutz
; STREET: 1220 Market Street, P.O. Box 2207
; CITY: Wilmington
; STATE: Delaware
; COUNTRY: U.S.A.
; ZIP: 19899
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/985,526
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/608,845
; FILING DATE: 16-JUL-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: McMorow Jr., Robert G
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (302) 658-9141
; TELEFAX: (302) 658-5613
; INFORMATION FOR SEQ ID NO: 36:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 185 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
US-08-985-526-36

Query Match      94.6%; Score 916; DB 3; Length 185;
Best Local Similarity 95.1%; Pred. No. 1.1e-105;
Matches 176; Conservative 5; Mismatches 2; Indels 2; Gaps 2;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRFLSSRLQDLYSI 60
Db 2 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFNNAR-VGLSGTFRFLSSRLQDLYSI 60

QY 61 VRRADRGSPVIVNLKDEVLSLSPSWDSLFSGSQGQLQPGARIFSDGRDVLHRHPAWPKSV 119
Db 61 VRRADRGSPVIVQNLDEVLSLSPSWDSLFSGSQGQLQPGARIFSDGRDVLHRHPAWPKSV 120

QY 120 WEGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 179
Db 121 WEGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLBQKAASCHNSYIVLCIENSEFMT 180
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Qy	180	TSESK	184
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Db	181	TSESR	185

RESULT 8
US-09-561-500-14
; Sequence 14, Application US/09561500
; Patent No. 6342219
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY COMPOSITIONS FOR SELECTIVELY INHIBITING VEGF
; FILE REFERENCE: 4001.002500
; CURRENT APPLICATION NUMBER: US/09/561,500
; CURRENT FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/131,432
; PRIOR FILING DATE: 1999-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 182
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
; OTHER INFORMATION: PEPTIDE
US-09-561-500-14

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Query Match      86.8%; Score 840; DB 4; Length 182;
- Best Local Similarity 85.6%; Pred. NO. 2.9e-96;
Matches 155; Conservative 15; Mismatches 11; Indels 0; Gaps 0;

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QY	1	HTHQDFQPVHLHVALNTPLSGGMRGIRGADFCFQQARAVGLSGTFRAFLSSRLQDLYSI	60
Db	1	HSHRDQFPVHLHVALNSPLSGMRCIRGADFCFQQARAVGLAGTFRFAFLSSRLQDLYSI	60
QY	61	VRRADRGSPPIVNKDEVLPSPWDSLFGSQOQLPGARIFSFDGRDVLRHHPWPQKSVM	120
Db	61	VRRADRRAVPINVNKDELLFPSWEALFGSEGPKPGARIFSFDGKDVLRHPTWPQKSVM	120
QY	121	HGSDPSGRRLMESYCETWRTEITTCATGOASSLLSGRLLLEOKAASCHNSYIVLCIENSFMT	180
Db	121	HGSDPNGRRLTESYCETWRTEAPSATGOASSLLGGRLLGSAASCHHAYIVLCIENSFMT	180
QY	181	S	181
Db	181	A	181

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RESULT 9
US-09-561-108-14
; Sequence 14, Application US/09561108
; Patent No. 6342221
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY CONJUGATE COMPOSITIONS FOR SELECTIVELY INHIBITING VEGF
; FILE REFERENCE: 4001.002584
; CURRENT APPLICATION NUMBER: US/09/561,108
; CURRENT FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/131,432
; PRIOR FILING DATE: 1999-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 182
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
; OTHER INFORMATION: PEPTIDE
US-09-561-108-14

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Query Match 86.8%; Score 840; DB 4; Length 182;
Best Local Similarity 85.6%; Pred. NO. 2.9e-96;
Matches 155; Conservative 15; Mismatches 11; Indels 0; Gaps 0;

QY	1	HTHQDFQPVHLHVALNTPLSGGMRGIRGADFCFQQARAVGLSGTFRAFLSSRLQLYSI	60
Dd	1	HSHRDFQPVHLHVALNTPLSGGMRGIRGADFCFQQARAVGLAGTFRAFLSSRLQLYSI	60
QY	61	VRRADRCSPVIVNLKDEVLPSPWDSLFSGSQGOLPGGARIFSFDRDVLRHPPAWPOKSVM	120
Dd	61	VRRADRRAVPVIVNLKDELLFPSWEALFSGSEGPLKPGARIFSFDGKDVLRHPTWPQKSVM	120
QY	121	HGSDPSGRRLMESYCETWRTEATTGATQAASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT	180
Dd	121	HGSDPNGRRLTESYCETWRTEAPSATGQAASSLLGGRLLGQSAASCHHAYIVLCIENSFMT	180
QY	181	S 181	
Dd	181	A 181	

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RESULT 10
US-09-315-689-3
; Sequence 3, Application US/09315689
; Patent No. 6346510
; GENERAL INFORMATION:
; APPLICANT: Folkman, Judah
; APPLICANT: O'Reilly, Michael
; TITLE OF INVENTION: Therapeutic Antiangiogenic Endostatin Compositions
; FILE REFERENCE: 05213-0229
; CURRENT APPLICATION NUMBER: US/09/315,689
; CURRENT FILING DATE: 1999-05-20
; NUMBER OF SEQ ID NOS: 6
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 3
; LENGTH: 182
; TYPE: PRT
; ORGANISM: Homo sapiens
US-09-315-689-3

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Query Match	86.8%;	Score 840;	DB 4;	Length 182;
Best Local Similarity	85.6%;	Pred. No. 2.9e-96;		
Matches 155:	Conservative	15;	Mismatches 11;	Indels 0;
				Gaps 0

QY	1	HTHQDFQPVHLHVALNTPLSGMRGIRGADFQCFQQAARAVCLSGTFRAFLSSRLQDLYSI	60
Db	1	HSHRDFQPVHLHVALNTPLSGMRGIRGADFQCFQQAARAVCLAGTFRAFLSSRLQDLYSI	60
QY	61	VRRADGSPVPIVNLKDEVLSFSDSLFSGSQGQLQPGARIFSFDRDVLRHHPAWPKSVW	120
Db	61	VRRADRAAVPIVNLKDELLFPWEALFSGSEGPLKPGARIFSFDKDVLRHPTWPKSVW	120
QY	121	HGSDPSSGRRLMESYCYETWRTTGTATGQASSLLSGRLLQKAASCHNSYIVLCIENSFMT	180
Db	121	HGSDPNGRRLTESYCYETWRTTEAPSATGQASSLLGGRLLQQAASCHHAYIVLCIENSFMT	180
QY	181	S	181
Db	181	A	181

RESULT 11
US-09-561-526-14
; Sequence 14, Application US/09561526
; Patent No. 6416758
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY CONJUGATE
; FILE REFERENCE: 4001.002586
; CURRENT APPLICATION NUMBER: US/09/561-526-14
; CURRENT FILING DATE: 2000-04-28

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 31, 2004, 19:58:01 ; Search time 125 Seconds
(without alignments)
463.110 Million cell updates/sec

Title: US-09-589-777C-2
Perfect score: 968
Sequence: 1 HTHQDFQPVHLVALNTPLS.....CHNSYIVLCIENSFMTSFSK 184

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1297172 seqs, 314612898 residues

Total number of hits satisfying chosen parameters: 1297172

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Published Applications_AA:*

1:	/cgn2_6/ptodata/2/pubpaa/US07_PUBCOMB.pep:*
2:	/cgn2_6/ptodata/2/pubpaa/PCT_NEW_PUB.pep:*
3:	/cgn2_6/ptodata/2/pubpaa/US06_NEW_PUB.pep:*
4:	/cgn2_6/ptodata/2/pubpaa/US06_PUBCOMB.pep:*
5:	/cgn2_6/ptodata/2/pubpaa/PCTUS_PUBCOMB.pep:*
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7:	/cgn2_6/ptodata/2/pubpaa/US08_PUBCOMB.pep:*
8:	/cgn2_6/ptodata/2/pubpaa/US09A_PUBCOMB.pep:*
9:	/cgn2_6/ptodata/2/pubpaa/US09B_PUBCOMB.pep:*
10:	/cgn2_6/ptodata/2/pubpaa/US09C_PUBCOMB.pep:*
11:	/cgn2_6/ptodata/2/pubpaa/US09_NEW_PUB.pep:*
12:	/cgn2_6/ptodata/2/pubpaa/US10A_PUBCOMB.pep:*
13:	/cgn2_6/ptodata/2/pubpaa/US10B_PUBCOMB.pep:*
14:	/cgn2_6/ptodata/2/pubpaa/US10C_PUBCOMB.pep:*
15:	/cgn2_6/ptodata/2/pubpaa/US10_NEW_PUB.pep:*
16:	/cgn2_6/ptodata/2/pubpaa/US60_NEW_PUB.pep:*
17:	/cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pep:*
18:	/cgn2_6/ptodata/2/pubpaa/US60_PUBCOMB.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	968	100.0	191	9	US-09-998-831-13
2	968	100.0	191	14	US-10-373-561-13
3	968	100.0	207	13	US-10-080-797-3
4	965	99.7	184	14	US-10-131-241-46
5	965	99.7	184	14	US-10-292-418-18
6	965	99.7	207	12	US-09-373-938-2
7	965	99.7	207	14	US-10-422-934-71
8	916	94.6	185	13	US-10-036-869-36
9	841	86.9	184	14	US-10-131-241-49
10	841	86.9	184	14	US-10-292-418-35
11	840	86.8	181	14	US-10-131-241-55
12	840	86.8	182	9	US-09-998-831-14
13	840	86.8	182	14	US-10-131-241-54
14	840	86.8	182	14	US-10-042-347-3
15	840	86.8	182	14	US-10-373-561-14

16	840	86.8	183	9	US-09-873-676-2	Sequence 2, Appli
17	840	86.8	183	12	US-09-978-531-1	Sequence 1, Appli
18	840	86.8	183	12	US-10-135-872B-11	Sequence 11, Appli
19	840	86.8	183	13	US-10-080-797-1	Sequence 1, Appli
20	840	86.8	183	14	US-10-131-241-52	Sequence 52, Appli
21	840	86.8	183	14	US-10-292-418-4	Sequence 4, Appli
22	840	86.8	183	16	US-10-607-501-2	Sequence 2, Appli
23	840	86.8	184	16	US-10-449-609-4	Sequence 4, Appli
24	840	86.8	208	12	US-09-373-938-5	Sequence 5, Appli
25	840	86.8	385	16	US-10-449-609-6	Sequence 6, Appli
26	840	86.8	682	15	US-10-264-049-3010	Sequence 3010, Ap
27	840	86.8	684	10	US-09-961-403-5	Sequence 5, Appli
28	840	86.8	1516	14	US-10-060-036-166	Sequence 166, App
29	840	86.8	1516	15	US-10-431-642-3	Sequence 3, Appli
30	839	86.7	180	14	US-10-131-241-56	Sequence 56, Appli
31	831	85.8	184	10	US-09-938-391-4	Sequence 4, Appli
32	831	85.8	230	10	US-09-938-391-2	Sequence 2, Appli
33	828	85.5	180	14	US-10-131-241-47	Sequence 47, Appli
34	823	85.0	184	12	US-10-210-172-162	Sequence 162, App
35	822	84.9	178	14	US-10-131-241-60	Sequence 60, Appli
36	822	84.9	178	14	US-10-042-347-5	Sequence 5, Appli
37	822	84.9	179	14	US-10-131-241-57	Sequence 57, Appli
38	791	81.7	171	12	US-09-978-531-3	Sequence 3, Appli
39	621	64.2	139	12	US-09-978-531-12	Sequence 12, Appli
40	583	60.2	160	12	US-10-210-172-164	Sequence 164, App
41	574	59.3	1279	12	US-10-087-192-1455	Sequence 1455, Ap
42	498	51.4	160	12	US-10-210-172-166	Sequence 166, App
43	294	30.4	63	9	US-09-822-540A-1	Sequence 1, Appli
44	294	30.4	63	12	US-09-978-531-5	Sequence 5, Appli
45	294	30.4	63	12	US-09-978-531-11	Sequence 11, Appli

ALIGNMENTS

RESULT 1
US-09-998-831-13
; Sequence 13, Application US/09998831
; Patent No. US20020119153A1
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY CONJUGATE COMPOSITIONS FOR SELECTIVELY
; TITLE OF INVENTION: INHIBITING VEGF
; FILE REFERENCE: 4001.002584
; CURRENT APPLICATION NUMBER: US/09/998,831
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: 09/561,108
; PRIOR FILING DATE: 2000-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
; LENGTH: 191
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-09-998-831-13

Query Match 100.0%; Score 968; DB 9; Length 191;
Best Local Similarity 100.0%; Pred. No. 1.1e-100;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY	1	HTHQDFQPVHLVALNTPLSGMRGIRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI	60
DB	8	HTHQDFQPVHLVALNTPLSGMRGIRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI	67
QY	61	VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPQKSVW	120
DB	68	VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPQKSVW	127
QY	121	HGSDPSGRRLMESYCYETWRTTTCATGQASSLLSGILLEQKAASCHNSYIVLCIENSFMT	180

Db 128 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 187

QY 181 SFSK 184
Db 188 SFSK 191

RESULT 2

US-10-373-561-13
; Sequence 13, Application US/10373561
; Publication No. US20030175276A1
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY METHODS FOR SELECTIVELY INHIBITING VEGF
; FILE REFERENCE: 4001.002582
; CURRENT APPLICATION NUMBER: US/10/373,561
; CURRENT FILING DATE: 2003-02-24
; PRIOR APPLICATION NUMBER: US/09/561,499
; PRIOR FILING DATE: 2000-04-28
; PRIOR APPLICATION NUMBER: 60/131,432
; PRIOR FILING DATE: 1999-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 13
; LENGTH: 191
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
US-10-373-561-13

Query Match 100.0%; Score 968; DB 14; Length 191;
Best Local Similarity 100.0%; Pred. No. 1.1e-100;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 67
QY 61 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 120
Db 68 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 127
QY 121 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
Db 128 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 187

QY 181 SFSK 184
Db 188 SFSK 191

RESULT 3

US-10-080-797-3
; Sequence 3, Application US/10080797
; Publication No. US20020183253A1
; GENERAL INFORMATION:
; APPLICANT: Campochiaro, Peter A.
; APPLICANT: Dixon, Katharine H.
; APPLICANT: Brazzell, Romulus K.
; TITLE OF INVENTION: METHOD FOR TREATING OCULAR
; TITLE OF INVENTION: NEOVASCULARIZATION
; FILE REFERENCE: 4-31881A
; CURRENT APPLICATION NUMBER: US/10/080,797
; CURRENT FILING DATE: 2002-02-21
; NUMBER OF SEQ ID NOS: 21
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 3
; LENGTH: 207
; TYPE: PRT
; ORGANISM: Mouse
US-10-080-797-3

Query Match 100.0%; Score 968; DB 13; Length 207;
Best Local Similarity 100.0%; Pred. No. 1.3e-100;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 24 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 83
QY 61 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 120
Db 84 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 143
QY 121 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
Db 144 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 203
QY 181 SFSK 184
Db 204 SFSK 207

RESULT 4

US-10-131-241-46
; Sequence 46, Application US/10131241
; Publication No. US20030012792A1
; GENERAL INFORMATION:
; APPLICANT: Holaday, John W.
; APPLICANT: Fortier, Anne H.
; TITLE OF INVENTION: Compositions and Methods for Inhibiting Endothelial Cell Proliferation
; TITLE OF INVENTION: and Regulating Angiogenesis Using Cancer Markers
; FILE REFERENCE: 05213-0344 43170-271565
; CURRENT APPLICATION NUMBER: US/10/131,241
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: US 09/413,049
; PRIOR FILING DATE: 1999-10-06
; PRIOR APPLICATION NUMBER: US 09/316,802
; PRIOR FILING DATE: 1999-05-21
; PRIOR APPLICATION NUMBER: US 60/086,586
; PRIOR FILING DATE: 1998-05-22
; NUMBER OF SEQ ID NOS: 65
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 46
; LENGTH: 184
; TYPE: PRT
; ORGANISM: Murinae sp.
US-10-131-241-46

Query Match 99.7%; Score 965; DB 14; Length 184;
Best Local Similarity 99.5%; Pred. No. 2.4e-100;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HTHQDFQPVHLVALNTPLSGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 120
Db 61 VRRADRGSVPIVNLKDEVLSFSGQQLPGARIFSFDFGRDVLRLHPAWPKSVW 120
QY 121 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
Db 121 HGS DPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
QY 181 SFSK 184
Db 181 SFSK 184

RESULT 5

US-10-292-418-18
; Sequence 18, Application US/10292418
; Publication No. US20030139365A1

GENERAL INFORMATION:
APPLICANT: Lo, Kin-Ming
APPLICANT: Li, Yue
APPLICANT: Gillies, Stephen D
TITLE OF INVENTION: Expression and Export of Angiogenesis Inhibitors as
TITLE OF INVENTION: Immunofusins
FILE REFERENCE: LEX-006C1
CURRENT APPLICATION NUMBER: US/10/292,418
CURRENT FILING DATE: 2002-11-12
PRIOR APPLICATION NUMBER: 09/383,315
PRIOR FILING DATE: 1999-08-25
PRIOR APPLICATION NUMBER: US 60/097,883
PRIOR FILING DATE: 1998-08-25
NUMBER OF SEQ ID NOS: 54
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 18
LENGTH: 184
TYPE: PRT
ORGANISM: Mus musculus
US-10-292-418-18

Query Match 99.7%; Score 965; DB 14; Length 184;
Best Local Similarity 99.5%; Pred. No. 2.4e-100;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQOQARAVGLSGTFRFLSSRLQDLYSI 60
DB 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQOQARAVGLSGTFRFLSSRLQDLYSI 60

QY 61 VRRADRGSPVIVNLKDEVLSFSGSQGQLOPGARIFSFDRDVLHRHPAWPKSVW 120
DB 61 VRRADRGSPVIVNLKDEVLSFSGSQGQLOPGARIFSFDRDVLHRHPAWPKSVW 120

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
DB 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

QY 181 SFSK 184
DB 181 SFSK 184

RESULT 6
US-09-373-938-2
Sequence 2, Application US/09373938
Publication No. US20020115202A1
GENERAL INFORMATION:
APPLICANT: Hallenbeck, Paul
APPLICANT: Chen, Cheauyun Theresa
TITLE OF INVENTION: ADENOVIRAL VECTORS INCLUDING DNA SEQUENCES ENCODING ANGIOGENIC IN
FILE REFERENCE: 4-30899P1
CURRENT APPLICATION NUMBER: US/09/373,938
CURRENT FILING DATE: 1999-08-13
NUMBER OF SEQ ID NOS: 17
SOFTWARE: PatentIn version 3.1
SEQ ID NO 2
LENGTH: 207
TYPE: PRT
ORGANISM: Mus musculus
US-09-373-938-2

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
DB 144 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 203

QY 181 SFSK 184
DB 204 SFSK 207

RESULT 7
US-10-422-934-71
Sequence 71, Application US/10422934
Publication No. US20030186841A1
GENERAL INFORMATION:
APPLICANT: Barbas, Carlos F., III
APPLICANT: Kadan, Michael
APPLICANT: Beerli, Roger
TITLE OF INVENTION: LIGAND ACTIVATED TRANSCRIPTIONAL REGULATOR PROTEINS
FILE REFERENCE: 22908-1227C
CURRENT APPLICATION NUMBER: US/10/422,934
CURRENT FILING DATE: 2003-04-23
PRIOR APPLICATION NUMBER: 09/586,625
PRIOR FILING DATE: 2000-06-02
PRIOR APPLICATION NUMBER: 09/433,042
PRIOR FILING DATE: 1999-10-25
NUMBER OF SEQ ID NOS: 92
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 71
LENGTH: 207
TYPE: PRT
ORGANISM: Muridae
US-10-422-934-71

Query Match 99.7%; Score 965; DB 14; Length 207;
Best Local Similarity 99.5%; Pred. No. 2.8e-100;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQOQARAVGLSGTFRFLSSRLQDLYSI 60
DB 24 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQOQARAVGLSGTFRFLSSRLQDLYSI 83

QY 61 VRRADRGSPVIVNLKDEVLSFSGSQGQLOPGARIFSFDRDVLHRHPAWPKSVW 120
DB 84 VRRADRGSPVIVNLKDEVLSFSGSQGQLOPGARIFSFDRDVLHRHPAWPKSVW 143

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
DB 144 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 203

QY 181 SFSK 184
DB 204 SFSK 207

RESULT 8
US-10-036-869-36
Sequence 36, Application US/10036869
Publication No. US20020151516A1
GENERAL INFORMATION:
APPLICANT: Mixson, James A
TITLE OF INVENTION: CARRIER:DNA COMPLEXES CONTAINING DNA
ENCODING ANTI-ANGIOGENIC PEPTIDES AND THEIR USE IN GENE
THERAPY
NUMBER OF SEQUENCES: 43
CORRESPONDENCE ADDRESS:
ADDRESSEE: Connolly, Bove, Lodge, & Hutz
STREET: 1220 Market Street, P.O. Box 2207
CITY: Wilmington
STATE: Delaware
COUNTRY: U.S.A.
ZIP: 19899
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/036,869
FILING DATE: 29-NO. US20020151516A1-2001
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/08/985,526
FILING DATE: <Unknown>
APPLICATION NUMBER: US 08/608,845
FILING DATE: 16-JUL-1996
ATTORNEY/AGENT INFORMATION:
NAME: McMorow Jr., Robert G
TELEPHONE: (302) 658-9141
TELEFAX: (302) 658-5613
INFORMATION FOR SEQ ID NO: 36:
SEQUENCE CHARACTERISTICS:
LENGTH: 185 amino acids
TYPE: amino acid
TOPOLOGY: linear
SEQUENCE DESCRIPTION: SEQ ID NO: 36:
US-10-036-869-36

Query Match 94.6%; Score 916; DB 13; Length 185;
Best Local Similarity 95.1%; Pred. No. 8.1e-95;
Matches 176; Conservative 5; Mismatches 2; Indels 2; Gaps 2;
QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
Db 2 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 119
Db 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 120
QY 120 WHGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 179
Db 121 WHGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 180
QY 180 TSFSK 184
Db 181 TSFSR 185

RESULT 9
US-10-131-241-49
Sequence 49, Application US/10131241
Publication No. US20030012792A1
GENERAL INFORMATION:
APPLICANT: Holaday, John W.
APPLICANT: Fortier, Anne H.
TITLE OF INVENTION: Compositions and Methods for Inhibiting Endothelial Cell Proliferation and Regulating Angiogenesis Using Cancer Markers
FILE REFERENCE: 05213-0344 43170-271565
CURRENT APPLICATION NUMBER: US/10/131,241
FILING DATE: 2002-07-22
PRIOR APPLICATION NUMBER: US 09/413,049
PRIOR FILING DATE: 1999-10-06
PRIOR APPLICATION NUMBER: US 09/316,802
PRIOR FILING DATE: 1999-05-21
PRIOR APPLICATION NUMBER: US 60/086,586
PRIOR FILING DATE: 1998-05-22
NUMBER OF SEQ ID NOS: 65
SOFTWARE: PatentIn version 3.1
SEQ ID NO 49
LENGTH: 184
TYPE: PRT
ORGANISM: Canine sp.
US-10-131-241-49
Query Match 86.9%; Score 841; DB 14; Length 184;
US-10-131-241-55
Sequence 55, Application US/10131241
Publication No. US20030012792A1
GENERAL INFORMATION:
APPLICANT: Holaday, John W.
APPLICANT: Fortier, Anne H.

Best Local Similarity 84.2%; Pred. No. 2.3e-86;
Matches 155; Conservative 17; Mismatches 12; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 120
Db 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 120
QY 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 180
Db 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 180
QY 181 SFSK 184
Db 181 SFSK 184

RESULT 10
US-10-292-418-35
Sequence 35, Application US/10292418
Publication No. US20030139365A1
GENERAL INFORMATION:
APPLICANT: Lo, Kin-Ming
APPLICANT: Li, Yue
APPLICANT: Gillies, Stephen D
TITLE OF INVENTION: Expression and Export of Angiogenesis Inhibitors as
FILE REFERENCE: LEX-006C1
CURRENT APPLICATION NUMBER: US/10/292,418
FILING DATE: 2002-11-12
PRIOR APPLICATION NUMBER: 09/383,315
PRIOR FILING DATE: 1999-08-25
PRIOR APPLICATION NUMBER: US 60/097,883
PRIOR FILING DATE: 1998-08-25
NUMBER OF SEQ ID NOS: 54
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 35
LENGTH: 184
TYPE: PRT
ORGANISM: Canis familiaris
US-10-292-418-35

Query Match 86.9%; Score 841; DB 14; Length 184;
Best Local Similarity 84.2%; Pred. No. 2.3e-86;
Matches 155; Conservative 17; Mismatches 12; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HTHQDFQPVHLVALNTPLSGMGRGADFCQFQARAVGLSGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 120
Db 61 VRRADRGSPVIVNLKDEVLSFSGQQLQPGARIFSFDRDVLHRHPAPQKSV 120
QY 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 180
Db 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGLRLEQKAAACHNSYIVLCIENSFM 180
QY 181 SFSK 184
Db 181 SFSK 184

RESULT 11
US-10-131-241-55
Sequence 55, Application US/10131241
Publication No. US20030012792A1
GENERAL INFORMATION:
APPLICANT: Holaday, John W.
APPLICANT: Fortier, Anne H.

;; TITLE OF INVENTION: Compositions and Methods for Inhibiting Endothelial Cell Proliferation
;; TITLE OF INVENTION: and Regulating Angiogenesis Using Cancer Markers
;; FILE REFERENCE: 05213-0344 43170-271565
;; CURRENT APPLICATION NUMBER: US/10/131,241
;; CURRENT FILING DATE: 2002-07-22
;; PRIOR APPLICATION NUMBER: US 09/413,049
;; PRIOR FILING DATE: 1999-10-06
;; PRIOR APPLICATION NUMBER: US 09/316,802
;; PRIOR FILING DATE: 1999-05-21
;; PRIOR APPLICATION NUMBER: US 60/086,586
;; PRIOR FILING DATE: 1998-05-22
;; NUMBER OF SEQ ID NOS: 65
;; SOFTWARE: PatentIn version 3.1
;; SEQ ID NO 55
;; LENGTH: 181
;; TYPE: PRT
;; ORGANISM: Homo sapiens
US-10-131-241-55

Query Match 86.8%; Score 840; DB 14; Length 181;
Best Local Similarity 85.6%; Pred. No. 2.9e-86;
Matches 155; Conservative 15; Mismatches 11; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HSHRDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLAGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSFSGQWDSLSPSWDLSFSGQQLQPGARIFSDGDRDVLHRHPAWPKSVW 120
Db 61 VRRADRAAVPIVNLKDELLFPSEALFSGSEGPKLPGARIFSDGDKDVLHRHTWPQKSVW 120
QY 121 HGSDPSGRRRLMESYCETWRTTETGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 121 HGSDPNGRRLTESYCETWRTTEAPSATGQASSLLGGRLLGQSAASCHHAYIVLCIENSFMT 180
QY 181 S 181
Db 181 A 181

RESULT 12
US-09-998-831-14
; Sequence 14, Application US/09998831
; Patent No. US20020119153A1
; GENERAL INFORMATION:
; APPLICANT: Philip E. Thorpe
; APPLICANT: Rolf A. Brekken
; TITLE OF INVENTION: ANTIBODY CONJUGATE COMPOSITIONS FOR SELECTIVELY
; TITLE OF INVENTION: INHIBITING VEGF
; FILE REFERENCE: 4001.002584
; CURRENT APPLICATION NUMBER: US/09/998,831
; CURRENT FILING DATE: 2001-11-30
; PRIOR APPLICATION NUMBER: 09/561,108
; PRIOR FILING DATE: 2000-04-28
; NUMBER OF SEQ ID NOS: 44
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 14
; LENGTH: 182
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: SYNTHETIC
; OTHER INFORMATION: PEPTIDE
US-09-998-831-14

Query Match 86.8%; Score 840; DB 9; Length 182;
Best Local Similarity 85.6%; Pred. No. 2.9e-86;
Matches 155; Conservative 15; Mismatches 11; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HSHRDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLAGTFRFLSSRLQDLYSI 60

QY 61 VRRADRGSPVIVNLKDEVLSFSGQWDSLSPSWDLSFSGQQLQPGARIFSDGDRDVLHRHPAWPKSVW 120
Db 61 VRRADRAAVPIVNLKDELLFPSEALFSGSEGPKLPGARIFSDGDKDVLHRHTWPQKSVW 120
QY 121 HGSDPSGRRRLMESYCETWRTTETGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 121 HGSDPNGRRLTESYCETWRTTEAPSATGQASSLLGGRLLGQSAASCHHAYIVLCIENSFMT 180
QY 181 S 181
Db 181 A 181

RESULT 13
US-10-131-241-54
; Sequence 54, Application US/10131241
; Publication No. US20030012792A1
; GENERAL INFORMATION:
; APPLICANT: Holaday, John W.
; APPLICANT: Fortier, Anne H.
; TITLE OF INVENTION: Compositions and Methods for Inhibiting Endothelial Cell Proliferation
; TITLE OF INVENTION: and Regulating Angiogenesis Using Cancer Markers
; FILE REFERENCE: 05213-0344 43170-271565
; CURRENT APPLICATION NUMBER: US/10/131,241
; CURRENT FILING DATE: 2002-07-22
; PRIOR APPLICATION NUMBER: US 09/413,049
; PRIOR FILING DATE: 1999-10-06
; PRIOR APPLICATION NUMBER: US 09/316,802
; PRIOR FILING DATE: 1999-05-21
; PRIOR APPLICATION NUMBER: US 60/086,586
; PRIOR FILING DATE: 1998-05-22
; NUMBER OF SEQ ID NOS: 65
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 54
; LENGTH: 182
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-131-241-54

Query Match 86.8%; Score 840; DB 14; Length 182;
Best Local Similarity 85.6%; Pred. No. 2.9e-86;
Matches 155; Conservative 15; Mismatches 11; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HSHRDFQPVHLVALNTPLSGGMRGIRGADFCQFQQAARAVGLAGTFRFLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSFSGQWDSLSPSWDLSFSGQQLQPGARIFSDGDRDVLHRHPAWPKSVW 120
Db 61 VRRADRAAVPIVNLKDELLFPSEALFSGSEGPKLPGARIFSDGDKDVLHRHTWPQKSVW 120
QY 121 HGSDPSGRRRLMESYCETWRTTETGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 121 HGSDPNGRRLTESYCETWRTTEAPSATGQASSLLGGRLLGQSAASCHHAYIVLCIENSFMT 180
QY 181 S 181
Db 181 A 181

RESULT 14
US-10-042-347-3
; Sequence 3, Application US/10042347
; Publication No. US20030114370A1
; GENERAL INFORMATION:
; APPLICANT: O'Reilly, Michael S.
; APPLICANT: Folkman, M. Judah
; TITLE OF INVENTION: Nucleic Acid Molecules Encoding Endostatin Protein and Peptide Fragments
; TITLE OF INVENTION: Thereof
; FILE REFERENCE: 05213-0880 (43170-249874)
; CURRENT APPLICATION NUMBER: US/10/042,347
; CURRENT FILING DATE: 2002-01-11
; PRIOR APPLICATION NUMBER: US 09/315,689

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OM protein - protein search, using sw model

Run on: August 31, 2004, 19:42:20 ; Search time 120 Seconds
(without alignments)
433.240 Million cell updates/sec

Title: US-09-589-777C-2
Perfect score: 968
Sequence: 1 HTHQDFQPVLHLVALNTPLS.....CHNSYIVLCIENSEMTSPSK 184

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1586107 seqs, 282547505 residues

Total number of hits satisfying chosen parameters: 1586107

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_29Jan04:*
1: Geneseqp1980s:*
2: Geneseqp1990s:*
3: Geneseqp2000s:*
4: Geneseqp2001s:*
5: Geneseqp2002s:*
6: Geneseqp2003as:*
7: Geneseqp2003bs:*
8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Query %			Match	Length	DB ID	Description
	Score	Match	Length				
1	968	100.0	184	4	AAB49380	Aab49380	Murine en
2	968	100.0	191	3	AAB28398	Aab28398	Murine en
3	968	100.0	191	5	AAU77950	Aau77950	Amino aci
4	968	100.0	207	5	ABB79902	Abb79902	Mouse end
5	965	99.7	184	2	AAU08689	Aay08689	Murine en
6	965	99.7	184	3	AAU70258	Aay70258	Murine an
7	965	99.7	184	5	ABG31793	Abg31793	Human end
8	965	99.7	207	4	AAE02031	Aae02031	Murine en
9	965	99.7	207	4	AAU71930	Aab71930	Murine en
10	965	99.7	218	2	AAU08691	Aay08691	Murine ge
11	965	99.7	580	2	AAU08692	Aay08692	Murine ge
12	965	99.7	1288	2	AAU26328	Aaw26328	Mouse alp
13	963	99.5	184	2	AAU18409	Aay18409	Endostati
14	960	99.2	1288	2	AAU92297	Aaw92297	Mouse alp
15	950	98.1	684	2	AAU25114	Aay25114	Mouse alp
16	946	97.7	183	5	AAU49504	Aam49504	Mouse end
17	916	94.6	185	2	AAU06197	Aay06197	Anti-angi
18	855	88.3	184	5	ABG31794	Abg31794	Murine en
19	841	86.9	184	3	AAU70265	Aay70265	Canine an
20	840	86.8	181	4	AAU00898	Aau00898	Human End
21	840	86.8	182	3	AAU59622	Aay59622	Human end
22	840	86.8	182	3	AAU94323	Aay94323	Human end
23	840	86.8	182	3	AAB28399	Aab28399	Human end
24	840	86.8	182	4	AAU00897	Aau00897	Human End
25	840	86.8	182	5	AAU77951	Aau77951	Amino aci

26	840	86.8	183	2	AAU02113	Aay02113	SEQ ID 76
27	840	86.8	183	2	AAU08693	Aay08693	Human end
28	840	86.8	183	3	AAU70252	Aay70252	Human ang
29	840	86.8	183	3	AAU90771	Aay90771	Human ang
30	840	86.8	183	3	AAB16451	Aab16451	Human end
31	840	86.8	183	3	AAB30493	Aab30493	Amino aci
32	840	86.8	183	4	AAB49379	Aab49379	Human end
33	840	86.8	183	4	AAU00896	Aau00896	Human End
34	840	86.8	183	5	ABB79901	Abb79901	Human end
35	840	86.8	183	5	AAM49503	Aam49503	Human end
36	840	86.8	183	5	AAM48895	Aam48895	Human end
37	840	86.8	183	5	AAU97132	Aau97132	Human end
38	840	86.8	183	6	AAG79753	Aag79753	Human end
39	840	86.8	195	3	AAU90874	Aaw90874	Human HMW
40	840	86.8	216	3	AAB30495	Aab30495	Amino aci
41	840	86.8	275	5	AAU76689	Aau76689	Synthetic
42	840	86.8	310	5	AAU76688	Aau76688	Human col
43	840	86.8	513	5	ABG73586	Abg73586	Human End
44	840	86.8	682	5	ABP41878	Abp41878	Human ova
45	840	86.8	684	2	AAW26327	Aaw26327	Human alp

ALIGNMENTS

RESULT 1
AAB49380
ID AAB49380 standard; protein; 184 AA.
XX

AC AAB49380;

DT 02-MAR-2001 (first entry)

XX 02-MAR-2001 (first entry)

DE Murine endostatin SEQ ID NO: 4.

XX Endostatin; antiangiogenic; angiogenesis; human; mouse; chicken; cancer;

KW inflammation; angiogenesis-dependent disease.

XX Mus musculus.

XX WC2000067771-A1.

XX 16-NOV-2000.

XX 02-MAY-2000; 2000WO-US012063.

XX 06-MAY-1999; 99US-0132907P.

PR 14-JUL-1999; 99US-00353333.

XX (BURN-) BURNHAM INST.

Vuori K;

WPI; 2001-0409337/05.

N-PSDB; AAC88290.

Endostatin peptide comprising at least four endostatin amino acid residues are e.g. angiogenesis inhibitors for treating cancer and diabetic retinopathy.

Disclosure; Fig 1; 146pp; English.

The present invention provides endostatin peptides which can be used in the modulation of angiogenesis. This is useful in the treatment of cancers, inflammation, rheumatoid arthritis, chronic articular rheumatism, psoriasis, disorders associated with inopportune invasion of vessels such as diabetic retinopathy, neovascular glaucoma, retinopathy of prematurity, macular degeneration, corneal graft rejection, retrolental fibroplasia, rubeosis, capillary proliferation in atherosclerotic plaques and osteoporosis. Other angiogenesis-dependent diseases include Osler-Webber syndrome, myocardial angiogenesis, plaque neovascularisation, telangiectasia, haemophilic joints and wound granulation. In addition, the peptides can be used as birth control

CC agents
XX
SQ Sequence 184 AA;

Query Match 100.0%; Score 968; DB 4; Length 184;
Best Local Similarity 100.0%; Pred. No. 1.3e-107;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGIRGADFQCFQARAVGLSGTFRAFLLSSRLQDLYSI 60
DB 1 HTHQDFQPVHLVALNTPLSGMGRGIRGADFQCFQARAVGLSGTFRAFLLSSRLQDLYSI 60
QY 61 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120
DB 61 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120
QY 121 HGSDPSGRRLMESYCETWRTTETTGATGQASSLLSGLLEQKAASCHNSYIVLCIENSEFMT 180
DB 121 HGSDPSGRRLMESYCETWRTTETTGATGQASSLLSGLLEQKAASCHNSYIVLCIENSEFMT 180
QY 181 SFSK 184
DB 181 SFSK 184

RESULT 2

AAB28398
ID AAB28398 standard; protein; 191 AA.

XX AC AAB28398;
DT 19-FEB-2001 (first entry)
XX Murine endostatin.
XX Murine; endostatin; cytostatic; antiproliferative;
KW vascular endothelial growth factor; VEGF; antibody; VEGF2 receptor;
KW cancer; vascularised solid tumour.
XX
OS Mus sp.
XX WO200064946-A2.
XX
PD 02-NOV-2000.
XX
PF 28-APR-2000; 2000WO-US011367.
XX
PR 28-APR-1999; 99US-0131432P.
XX
PA (TEXA) UNIV TEXAS SYSTEM.
XX
PI Thorpe PE, Brekken RA;
XX WPI; 2000-687317/67.
DR N-PSDB; AAC67777.
XX
PT Immunogenic composition for the treatment and diagnosis of cancer
PT comprises an anti-VEGF (vascular endothelial growth factor) antibody
PT binding the same epitope as the monoclonal antibody ATCC PTA 1595.
XX
PS Example 10; Page 290-291; 298pp; English.
XX
CC The present invention relates to anti-Vascular Endothelial Growth Factor
CC (VEGF) antibodies that bind to the same epitope as the monoclonal
CC antibody ATCC PTA 1595 and which significantly inhibit VEGF binding to
CC the VEGF receptor VEGFR2, without inhibiting VEGF binding to the VEGF
CC receptor VEGFR1. The present sequence is murine endostatin. Endostatin
CC may be conjugated onto the anti-VEGF antibodies of the present invention.
CC The anti-VEGF antibodies of the present invention are useful for the
CC treatment and diagnosis of cancer, especially vascularised solid tumours
XX
SQ Sequence 191 AA;

Query Match 100.0%; Score 968; DB 3; Length 191;
Best Local Similarity 100.0%; Pred. No. 1.4e-107;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGIRGADFQCFQARAVGLSGTFRAFLLSSRLQDLYSI 60
DB 8 HTHQDFQPVHLVALNTPLSGMGRGIRGADFQCFQARAVGLSGTFRAFLLSSRLQDLYSI 67
QY 61 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120
DB 68 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 127
QY 121 HGSDPSGRRLMESYCETWRTTETTGATGQASSLLSGLLEQKAASCHNSYIVLCIENSEFMT 180
DB 128 HGSDPSGRRLMESYCETWRTTETTGATGQASSLLSGLLEQKAASCHNSYIVLCIENSEFMT 187
QY 181 SFSK 184
DB 188 SFSK 191

RESULT 3

AAU77950
ID AAU77950 standard; protein; 191 AA.

XX AC AAU77950;
DT 02-JUL-2002 (first entry)
XX Amino acid sequence for mouse endostatin.
DE Mouse; immunoconjugate; anti-vascular endothelial growth factor antibody;
XX anti-VEGF antibody; monoclonal antibody 2C3 ATCC PTA 1595; VEGF receptor;
KW VEGFR2; KDR/Flk-1; VEGFR1; Flt-1; angiogenesis; macular degeneration;
KW ocular neovascular disease; cancer; vascularised solid tumour; AIDS;
KW metastatic tumour; endothelial cell proliferation; inflammatory disorder;
KW atherosclerosis; diabetic retinopathy; corneal graft rejection;
KW acquired immune deficiency syndrome; infection; restenosis; fungal ulcer;
KW sickle cell anaemia; endometriosis; endostatin.
XX
OS Mus sp.
XX
PN AU200179401-A.
XX
PD 06-DEC-2001.
XX
PF 12-OCT-2001; 2001AU-00079401.
XX
PR 28-APR-2000; 2000AU-00048049.
XX
PA (TEXA) UNIV TEXAS SYSTEM.
XX
PI Thorpe PE, Brekken RA;
XX WPI; 2002-281368/33.
DR N-PSDB; ABK47719.
XX
PT Immunoconjugate compositions for treating cancer by inhibiting
PT angiogenesis and for delivering a diagnostic agent to tumor, comprises
PT anti-vascular endothelial growth factor antibody attached to a biological
PT agent.
XX
PS Example 10; Page 11-12 (Sequence listing); 299pp; English.
XX
CC The present invention relates to antibody-based compositions comprising
CC an immunoconjugate such as anti-vascular endothelial growth factor (VEGF)
CC antibody (Ab) (or its antigen-binding fragment), attached to a biological
CC agent, where the Ab binds to the same epitope as the monoclonal antibody
CC (Mab) 2C3 ATCC PTA 1595, and significantly inhibits VEGF binding to the
CC VEGF receptor VEGFR2 (KDR/Flk-1) without inhibiting VEGF binding to the
CC VEGF receptor VEGFR1 (Flt-1). The compositions of the invention are
CC useful in therapy, and diagnosis, for inhibiting angiogenesis in an
CC animal having ocular neovascular disease or macular degeneration, and for

CC delivering a biological agent to a vascularised tumour. The compositions
CC can also be used for treating cancer and subjects at risk of developing,
CC a vascularised solid tumour, a metastatic tumour or metastases from a
CC primary tumour. The composition is useful for specifically inhibiting
CC VEGF-induced endothelial cell proliferation, without significantly
CC inhibiting VEGF-induced macrophage, osteoclast or chondroclast function.
CC The compositions can be used for treating various diseases such as
CC inflammatory disorders, atherosclerosis, diabetic retinopathy,
CC restenosis, acquired immune deficiency syndrome (AIDS), blood borne
CC tumours, corneal graft rejection, Crohn's disease, fungal ulcers,
CC infections, sickle cell anaemia, and endometriosis. The present sequence
CC represents mouse endostatin. Endostatin may be attached or functionally
CC associated with anti-VEGF antibodies
XX
SQ Sequence 191 AA;

Query Match 100.0%; Score 968; DB 5; Length 191;
Best Local Similarity 100.0%; Pred. No. 1.4e-107;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 8 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQQAQAVGLSGTFRFLSSRLQDLYSI 67
Qy 61 VRRADRGSPVPIVNLKDEVLSFSGQQLQPGARIFSFDRDVLRRHPAWPKSVW 120
Db 68 VRRADRGSPVPIVNLKDEVLSFSGQQLQPGARIFSFDRDVLRRHPAWPKSVW 127
Qy 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 128 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 187
Qy 181 SFSK 184
Db 188 SFSK 191

RESULT 4
ABB79902
ID ABB79902 standard; protein; 207 AA.

XX AC ABB79902;
XX 05-DEC-2002 (first entry)
DT Mouse endostatin.
XX Endostatin; mouse; ophthalmological; ocular neovascularisation;
KW choroidal neovascularisation; gene therapy.
XX Mus musculus.
XX Key Location/Qualifiers
FH Misc-difference 117
FT /note= "encoded by GTG"

XX WO200267971-A2.
XX 06-SEP-2002.
XX 21-FEB-2002; 2002WO-US005336.
XX 22-FEB-2001; 2001US-0270787P.
PR 04-APR-2001; 2001US-0281296P.
XX (NOVS) NOVARTIS AG.
XX Brazzell RK, Campochiaro PA, Dixon KH;
XX WPI; 2002-698636/75.
DR N-PSDB; ABQ81194.

PT Treating or preventing choroidal neovascularization comprises increasing

PT the amount of endostatin in ocular tissues of afflicted individuals to a
PT choroidal neovascularization inhibiting level.

PS Disclosure; Page 40; 44pp; English.

XX The present sequence is the protein sequence of murine endostatin plus
CC the murine Ig kappa leader sequence. A claimed method for the treatment
CC of ocular neovascularisation, especially choroidal neovascularisation,
CC involves increasing the level of endostatin in ocular tissue, especially
CC the level of human endostatin (see ABB79901), its fragment, derivative or
CC variant. The increase is effected by administering a viral vector,
CC especially an adenovirus, adeno-associated virus, a retrovirus or
CC lentivirus vector, comprising an endostatin-encoding nucleic acid. Cells
CC secreting endostatin may be encapsulated and implanted within an
CC individual. The method is used when ocular neovascularisation is caused
CC by histoplasmosis, pathological myopia, angiod streaks, anterior
CC ischaemic optic neuropathy, bacterial endocarditis, Best's disease,
CC birdshot retinochoroidopathy, choroidal haemangioma, choroidal naevi,
CC choroidal nonperfusion, choroidal osteomas, choroidal rupture,
CC choroideraemia, chronic retinal detachment, coloboma of the retina,
CC Drusen, endogenous Candida endophthalmitis, extracapsillary hamartoma of
CC the retinal pigmented epithelium, fundus flavimaculatus, idiopathic,
CC macular hole, malignant melanoma, membranoproliferative glomerulonephritis
CC (type II), metallic intraocular foreign body, morning glory disc
CC syndrome, multiple evanescent white-dot syndrome, neovascularisation of
CC ora serrata, operating microscope burn, optic nerve head pits,
CC photocoagulation, punctate inner choroidopathy, rubella, sarcoidosis,
CC serpiginous or Geographic choroiditis, subretinal fluid drainage, tiled
CC disc syndrome, Toxoplasma retinochoroiditis, tuberculosis, Vogt-Koyanagi-
CC Harada syndrome, diabetic retinopathy, non-diabetic retinopathy, brain
CC vein occlusion, central retinal vein occlusion, retinopathy in premature
CC infants, rubosis iridis, neovascular glaucoma, perifoveal
CC telangiectasis, sickle cell retinopathy, Sallé's disease, retinal
CC vasculitis, Von Hippel Lindau disease, radiation retinopathy, retinal
CC cryoinjury, retinitis pigmentosa, retinochoroidal coloboma, corneal
CC neovascularisation due to herpes simplex keratitis, corneal ulcers,
CC keratoplasty, pterygia and trauma (all claimed)

XX SQ Sequence 207 AA;

Query Match 100.0%; Score 968; DB 5; Length 207;
Best Local Similarity 100.0%; Pred. No. 1.6e-107;
Matches 184; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 24 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQQAQAVGLSGTFRFLSSRLQDLYSI 83
Qy 61 VRRADRGSPVPIVNLKDEVLSFSGQQLQPGARIFSFDRDVLRRHPAWPKSVW 120
Db 84 VRRADRGSPVPIVNLKDEVLSFSGQQLQPGARIFSFDRDVLRRHPAWPKSVW 143
Qy 121 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 144 HGSDPSGRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 203
Qy 181 SFSK 184
Db 204 SFSK 207

RESULT 5
AAY08689
ID AAY08689 standard; protein; 184 AA.

XX AC AAY08689;
XX 10-AUG-1999 (first entry)
DT Murine endostatin protein fragment.
XX Plasmagen; murine; angiotatin; endostatin; gene therapy; vector;
KW anti-angiogenic; attenuation; cytostatic; anti-diabetic; ophthalmology;

KW tumour growth; solid tumour; diabetic retinopathy; retina.

XX Mus sp.

OS WO9926480-A1.

XX 03-JUN-1999.

XX 20-NOV-1998; 98WO-US024950.

XX 20-NOV-1997; 97US-00975424.

XX (GENE-) GENETIX PHARM INC.

XX (MASI) MASSACHUSETTS INST TECHNOLOGY.

XX Leboulch P, Pawliuk RJ, Bachelot T;

XX WPI; 1999-357696/30.

XX N-PSDB; AAX77715.

XX Anti-angiogenic gene therapy vectors.

XX Disclosure; Fig 6; 83pp; English.

CC This invention describes a novel viral gene therapy vector comprising a nucleic acid molecule encoding an anti-angiogenic polypeptide chosen from human or murine angiostatin, human or murine endostatin and angiogenesis-inhibiting fusions and fragments, where the viral vector is sufficiently attenuated for use in human gene therapy. The products of the invention have anti-angiogenic, cytostatic, anti-diabetic and ophthalmological activity. The vector is used in gene therapy for inhibiting tumour growth in humans harbouring a solid tumour. The vector expresses an anti-angiogenic polypeptide. An additional use comprises treatment of diabetic retinopathy, where the anti-angiogenic polypeptide inhibits angiogenesis in the vicinity of the retina. The vector is administered to cells ex vivo and then administered to the patient

XX Sequence 184 AA;

Query Match 99.7%; Score 965; DB 2; Length 184;
Best Local Similarity 99.5%; Pred. No. 3e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFFRAFLSSRLQDLYSI 60

Db 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFFRAFLSSRLQDLYSI 60

QY 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPWPKSVW 120

Db 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPWPKSVW 120

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

Db 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

QY 181 SFSK 184

Db 181 SFSK 184

RESULT 6

AAY70258

ID AAY70258 standard; protein; 184 AA.

XX AAY70258;

XX 06-JUN-2000 (first entry)

XX Murine angiogenesis inhibitor, endostatin.

XX Murine; immunoglobulin Fc fragment; endostatin; immunofusin;
KW angiogenesis; inhibitor; cytostatic; antirheumatoid; antiarthritic;
KW antipsoriatic; antidiabetic; ophthalmological; immunosuppressant;

KW

KW vasotropic; vulnery; treatment; antiarteriosclerosis; tumour;

KW metastasis; atherosclerosis; psoriasis; rheumatoid arthritis;

KW ocular angiogenic disease; diabetic retinopathy; macular degeneration;

KW myocardial angiogenesis; plaque neovascularisation; telangiectasia;

KW wound granulation; keloid scar; gene therapy.

XX Mus musculus.

XX WO200011033-A2.

XX 02-MAR-2000.

XX 25-AUG-1999; 99WO-US019329.

XX 25-AUG-1998; 98US-0097883P.

XX (LEXI-) LEXINGEN PHARM CORP.

XX Lo K, Li Y, Gillies SD;

XX WPI; 2000-237616/20.

XX N-PSDB; AAZ51299.

PT Novel fusion protein of angiostatin or endostatin and an immunoglobulin FC region, useful for treating conditions mediated by angiogenesis, such as rheumatoid arthritis, tumors and macular degeneration.

XX Example 5; Page 48-49; 68pp; English.

CC The patent discloses a DNA molecule encoding a fusion protein comprising a signal sequence, an immunoglobulin Fc region, and an angiogenesis inhibitor selected from angiostatin, endostatin, a plasminogen fragment having angiostatin activity, a collagen XVIII fragment having endostatin activity, or combinations of them. The fusion protein (immunofusin) is used to inhibit angiogenesis and to treat diseases or conditions mediated by angiogenesis. Conditions that may be treated include solid tumours, blood born tumours, tumour metastasis, benign tumours including haemangiomas, acoustic neuromas, neurofibromas, trachomas and pyrogenic granulomas, rheumatoid arthritis, psoriasis, ocular angiogenic diseases e.g. diabetic retinopathy, retinopathy of prematurity, macular degeneration, corneal graft rejection, neovascular glaucoma, retrolental fibroplasia, rubeosis and Osler-Webber syndrome; myocardial angiogenesis, plaque neovascularisation, telangiectasia, haemophilic joints,

CC angioblastoma, wound granulation, and excessive or abnormal stimulation of endothelial cells, intestinal cells, atherosclerosis, sclerodermal and hypertrophic scars, i.e. keloid scars. The DNA constructs may be used in gene therapy. The present sequence is a murine endostatin used in the construction of immunofusin containing murine immunoglobulin Fc fragment

XX Sequence 184 AA;

Query Match 99.7%; Score 965; DB 3; Length 184;
Best Local Similarity 99.5%; Pred. No. 3e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFFRAFLSSRLQDLYSI 60

Db 1 HTHQDFQPVHLVALNTPLSGMGRGADFCFQQAQAVGLSGTFFRAFLSSRLQDLYSI 60

QY 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPWPKSVW 120

Db 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSFQDGRDVLRHHPAPWPKSVW 120

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

Db 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

QY 181 SFSK 184

Db 181 SFSK 184

RESULT 7

ABG31793
ID ABC31793 standard; protein; 184 AA.
XX
AC ABC31793;
XX
DT 05-DEC-2002 (first entry)
XX
DE Human endostatin polypeptide.
XX
KW Human; endostatin; tumour; cancer; metastasis; cytostatic;
KW antiangiogenic.
XX
OS Homo sapiens.
XX
PN WO200268457-A2.
XX
PD 06-SEP-2002.
XX
PF 27-FEB-2002; 2002WO-IT000119.
XX
PR 27-FEB-2001; 2001IT-MI000394.
XX
PA (UYMI-) UNIV MILANO.
XX
PI Chillemi F, Vicentinie LMT, Francescato P;
XX
DR WPI; 2002-698655/75.
XX
PT New peptide useful for the preparation of medicaments with antiangiogenic
PT activity that may be used in treating tumors or metastases, comprises a
PT sequence corresponding to fragments of human endostatin.
XX
PS Disclosure; Fig 1; 24pp; English.
XX
CC The invention relates to peptide comprising 20-50 amino acids with
CC sequences corresponding to the human endostatin polypeptide sequence, its
CC salt or non-toxic derivative. The peptides are useful in the preparation
CC of medicaments with antiangiogenic activity which may be useful in
CC treating tumours or metastases. This sequence represents a human
CC endostatin polypeptide
XX
SQ Sequence 184 AA;

Query Match 99.7%; Score 965; DB 5; Length 184;
Best Local Similarity 99.5%; Pred. No. 3e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRAFLSSRLQDLYSI 60
DB 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRAFLSSRLQDLYSI 60

QY 61 VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120
DB 61 VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120

QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
DB 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

QY 181 SFSK 184
DB 181 SFSK 184

RESULT 8
AAE02031
ID AAE02031 standard; protein; 207 AA.
XX
AC AAE02031;
XX
DT 31-JUL-2001 (first entry)
XX
DE Murine endostatin fused to N-terminal secretion signal.

XX
KW Murine; endostatin; fusion protein; nucleotide-binding domain; NBD;
KW ligand-binding domain; LBD; transcription regulating domain; TRD; cancer;
KW zinc finger protein; ZFP; ligand-activated transcriptional regulator;
KW gene regulation; gene therapy; cell proliferative disorder; psoriasis;
KW pemphigus vulgaris; Behcet's syndrome; lipid histiocytosis.
XX
OS Mus sp.
XX
PN WO200130843-A1.
XX
PD 03-MAY-2001.
XX
PF 23-OCT-2000; 2000WO-EP010430.
XX
PR 25-OCT-1999; 99US-00433042.
PR 02-JUN-2000; 2000US-00586625.
XX
PA (NOVS) NOVARTIS AG.
PA (SCRI) SCRIPPS RES INST.
XX
PI Barbas CF, Kadan M, Beerli R;
XX
DR WPI; 2001-308618/32.
DR N-PSDB; AAD06108.
XX
PT New fusion protein containing nucleotide-binding and ligand-binding
PT domains, useful e.g. in gene therapy of cancer, provides ligand-activated
PT control of gene expression.
XX
PS Example 19; Page 209; 218pp; English.
XX
CC The invention relates to fusion protein comprising a nucleotide-binding
CC domain (NBD), a ligand-binding domain (LBD) of an intracellular receptor
CC (ICR) and a transcription regulating domain (TRD). NBD is a polydactyl
CC zinc finger protein (ZFP), or a modular part of it, that interacts
CC specifically with a contiguous sequence of at least 3 nucleotides. The
CC fusion protein functions as a ligand-activated transcriptional regulator.
CC The fusion protein and the nucleic acid encoding it, are used to regulate
CC gene expression, particularly in gene therapy for treating malignant cell
CC proliferative diseases (e.g. colon cancer, prostate cancer, renal-cell
CC carcinoma) and non-malignant cell proliferative diseases (e.g. psoriasis,
CC pemphigus vulgaris, Behcet's syndrome and lipid histiocytosis). The
CC fusion protein and its DNA are also useful for treating diseases caused
CC by viruses in humans/plants, genetic and/or acquired diseases. The fusion
CC protein can be designed to target any selected gene (endogenous or
CC exogenous), and can be made to have different selectivity or specificity
CC for endogenous or exogenous ligands. The present sequence is murine
CC endostatin fused to an N-terminal secretion signal. The corresponding
CC cDNA sequence was used in the construction of left end shuttle plasmids
CC containing regulatable transgene cassettes for evaluation of Cys2-His2
CC Zinc finger DNA binding domain (DBD)-Oestrogen receptor (ER) LBD
CC regulators
XX
SQ Sequence 207 AA;

Query Match 99.7%; Score 965; DB 4; Length 207;
Best Local Similarity 99.5%; Pred. No. 3.6e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRAFLSSRLQDLYSI 60
DB 24 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAARAVGLSGTFRAFLSSRLQDLYSI 83

QY 61 VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 120
DB 84 VRRADRGSPVIVNLKDEVLSFSGSQQLQPGARIFSFDRDVLRLHPAWPQKSVW 143

QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
DB 144 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 203

QY 181 SFSK 184

Db 204 SFSK 207
|||||
RESULT 9
AAB71930
ID AAB71930 standard; protein; 207 AA.
XX
AC AAB71930;
XX
DT 10-MAY-2001 (first entry)
XX
DE Murine endostatin attached to Ig-kappa leader sequence.
XX
KW Mouse; endostatin; antitumour; cytostatic; antiarthritic; antipsoriatic;
KW antidiabetic; ophthalmological; gene therapy; angiogenic inhibitor;
KW adenoviral vector; diabetic retinopathy; cardiovascular disease;
KW arthritis; psoriasis; cerebral oedema; intravascular coagulopathy;
KW lymphoma; leukaemia; immunoglobulin; Ig; Ig-kappa.
XX
OS Mus sp.
XX
PN WO200112830-A1.
XX
PD 22-FEB-2001.
XX
PF 11-AUG-2000; 2000WO-EP007865.
XX
PR 13-AUG-1999; 99US-00373938.
XX
PA (NOVS) NOVARTIS AG.
PA (NOVS) NOVARTIS-ERFINDUNGEN VERN GES MBH.
XX
PI Hallenbeck PL, Chen CT;
XX
DR WPI; 2001-202871/20.
DR N-PSDB; AAF60336.
XX
PT Adenoviral vector for treating tumors and disorders associated with
PT angiogenesis, such as cancer, arthritis, and psoriasis, comprises a DNA
PT sequence encoding an angiogenic inhibitor, particularly endostatin.
XX
PS Example 1; Fig 1B; 59pp; English.
XX
CC The nucleotide sequence encoding this protein was used in the
CC construction of an adenoviral vector which includes a DNA sequence
CC encoding endostatin. The adenoviral vector is useful for expressing
CC endostatin in a mammalian cell such as an A549 or Hep3B cell. It is
CC useful for treating other diseases and disorders associated with
CC angiogenesis, such as neovascular diseases of the eye, including diabetic
CC retinopathy, cardiovascular disease, arthritis, psoriasis, cerebral
CC oedema and intravascular coagulopathy (Kasabach-Merritt syndrome). The
CC vector inhibits, prevents or destroys the growth of tumours by preventing
CC the formation of blood vessels in tumours, such as lymphoma and leukaemia
XX
SQ Sequence 207 AA;

Query Match 99.7%; Score 965; DB 4; Length 207;
Best Local Similarity 99.5%; Pred. No. 3.6e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 24 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 83

QY 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSDGRDVLHRHPAPQKSVW 120
Db 84 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSDGRDVLHRHPAPQKSVW 143

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
Db 144 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 203

QY 181 SFSK 184
|||||
Db 204 SFSK 207

RESULT 10
AAY08691
ID AAY08691 standard; protein; 218 AA.
XX
AC AAY08691;
XX
DT 10-AUG-1999 (first entry)
XX
DE Murine gene therapy peptide construct SP-Flag-Endo.
XX
KW Plasminogen; murine; angiotatin; endostatin; gene therapy; vector;
KW anti-angiogenic; attenuation; cytostatic; anti-diabetic; ophthalmology;
KW tumour growth; solid tumour; diabetic retinopathy; retina; construct.
XX
OS Mus sp.
OS Synthetic.
XX
PN WO9926480-A1.
XX
PD 03-JUN-1999.
XX
PF 20-NOV-1998; 98WO-US024950.
XX
PR 20-NOV-1997; 97US-00975424.
XX
PA (GENE-) GENETIX PHARM INC.
PA (NASI) MASSACHUSETTS INST TECHNOLOGY.
XX
PI Leboulch P, Pawliuk RJ, Bachelot T;
XX
DR WPI; 1999-357696/30.
DR N-PSDB; AAX77717.
XX
PT Anti-angiogenic gene therapy vectors.
XX
PS Example 1; Page 69; 83pp; English.
XX
CC This invention describes a novel viral gene therapy vector comprising a
CC nucleic acid molecule encoding an anti-angiogenic polypeptide chosen from
CC human or murine angiotatin, human or murine endostatin and angiogenesis-
CC inhibiting fusions and fragments, where the viral vector is sufficiently
CC attenuated for use in human gene therapy. The products of the invention
CC have anti-angiogenic, cytostatic, anti-diabetic and ophthalmological
CC activity. The vector is used in gene therapy for inhibiting tumour growth
CC in humans harbouring a solid tumour. The vector expresses an anti-
CC angiogenic polypeptide. An additional use comprises treatment of diabetic
CC retinopathy, where the anti-angiogenic polypeptide inhibits angiogenesis
CC in the vicinity of the retina. The vector is administered to cells ex
CC vivo and then administered to the patient
XX
SQ Sequence 218 AA;

Query Match 99.7%; Score 965; DB 2; Length 218;
Best Local Similarity 99.5%; Pred. No. 3.9e-107;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 60
Db 35 HTHQDFQPVHLVALNTPLSGGMRGIRGADFCFQQAQAVGLSGTFRFLSSRLQDLYSI 94

QY 61 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSDGRDVLHRHPAPQKSVW 120
Db 95 VRRADRGSVPIVNLKDEVLSFSGSQQLQPGARIFSDGRDVLHRHPAPQKSVW 154

QY 121 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 180
Db 155 HGSDPSGRRRLMESYCETWRTTGTATGQASSLLSGRLLEQKAASCHNSYIVLCIENSFMT 214

Qy	181	SFSK	184
Db	215	SFSK	218

RESULT 11
 AAY08692
 ID AAY08692 standard; protein; 580 AA.

OS	Mus sp.	
OS	Synthetic.	
XX		
PN	WO9926480-A1.	
XX		
PD	03-JUN-1999.	
XX		
XX	20-NOV-1998;	98WO-US024950.
XX		
PR	20-NOV-1997;	97US-00975424.
XX		
PA	(GENE-) GENETIX PHARM INC.	
PA	(MASI) MASSACHUSETTS INST TECHNOLOGY.	
XX		
PPI	Leboulch P, Pawliuk RJ, Bachelot T;	
XX		
DR	WPI; 1999-357696/30.	
DR	N-PSDB; AAX77718.	

Anti-angiogenic gene therapy vectors.
 Example 1; Page 72-74; 83pp; English.

This invention describes a novel viral gene therapy vector comprising a nucleic acid molecule encoding an anti-angiogenic polypeptide chosen from human or murine angiostatin, human or murine endostatin and angiogenesis-inhibiting fusions and fragments, where the viral vector is sufficiently attenuated for use in human gene therapy. The products of the invention have anti-angiogenic, cytostatic, anti-diabetic and ophthalmological activity. The vector is used in gene therapy for inhibiting tumour growth in humans harbouring a solid tumour. The vector expresses an anti-angiogenic polypeptide. An additional use comprises treatment of diabetic retinopathy, where the anti-angiogenic polypeptide inhibits angiogenesis in the vicinity of the retina. The vector is administered to cells *ex vivo* and then administered to the patient.

Sequence 580 AA;

```

Query Match      99.7%; Score 965; DB 2; Length 580;
Best Local Similarity 99.5%; Pred. NO. 1.7e-106;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

```

Qy 1 HTHDFFPVLHLVALNTPUSGGMRGIRGADFQCFOQARAVGLSGTFRAFLSRLQLYSI 60
 |||||
Dd 397 HTHDFFPVLHLVALNTPUSGGMRGIRGADFQCFOQARAVGLSGTFRAFLSRLQLYSI 456

	61	VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQGLQPGARIFSFDGRDVLRHFAWPQKSVM	120
Qy		:	
	457	VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQGQQVGPGARIFSFDGRDVLRHFAWPQKSVM	516
Dd		:	

Qy

121	HGSDPSGRRLMESYCETWRTTETTATGGQAASSLLSGRILLEKKAASCHNSYIVLCIENSFMT	180

Dd

517	HGSDPSGRRLMESYCETWRTTETTATGGQAASSLLSGRILLEKKAASCHNSYVLCIENSFMT	576
-----	--	-----

FT Peptide /label= GXYGX'Y' _motif
FT 512. .517
FT /label= GXYGX'Y' _motif
FT 518. .523
FT /label= GXYGX'Y' _motif
FT 524. .529
FT /label= GXYGX'Y' _motif
FT 530. .535
FT /label= GXYGX'Y' _motif
FT 536. .541
FT /label= GXYGX'Y' _motif
FT 542. .547
FT /label= GXYGX'Y' _motif
FT 548. .553
FT /label= GXYGX'Y' _motif
FT 580. .585
FT /label= GXYGX'Y' _motif
FT 586. .591
FT /label= GXYGX'Y' _motif
FT 592. .597
FT /label= GXYGX'Y' _motif
FT 598. .603
FT /label= GXYGX'Y' _motif
FT 604. .609
FT /label= GXYGX'Y' _motif
FT 610. .615
FT /label= GXYGX'Y' _motif
FT 616. .621
FT /label= GXYGX'Y' _motif
FT 622. .627
FT /label= GXYGX'Y' _motif
FT 628. .633
FT /label= GXYGX'Y' _motif
FT 634. .639
FT /label= GXYGX'Y' _motif
FT 640. .665
FT /label= GXYGX'Y' _motif
FT 657. .662
FT /label= GXYGX'Y' _motif
FT 677. .682
FT /label= GXYGX'Y' _motif
FT 683. .688
FT /label= GXYGX'Y' _motif
FT 689. .694
FT /label= GXYGX'Y' _motif
FT 695. .700
FT /label= GXYGX'Y' _motif
FT 707. .712
FT /label= GXYGX'Y' _motif
FT 713. .718
FT /label= GXYGX'Y' _motif
FT 735. .740
FT /label= GXYGX'Y' _motif
FT 741. .746
FT /label= GXYGX'Y' _motif
FT 747. .752
FT /label= GXYGX'Y' _motif
FT 759. .764
FT /label= GXYGX'Y' _motif
FT 765. .770
FT /label= GXYGX'Y' _motif
FT 771. .776
FT /label= GXYGX'Y' _motif
FT 787. .792
FT /label= GXYGX'Y' _motif
FT 793. .798
FT /label= GXYGX'Y' _motif
FT 799. .804
FT /label= GXYGX'Y' _motif
FT 815. .820
FT /label= GXYGX'Y' _motif
FT 821. .826
FT /label= GXYGX'Y' _motif

FT Peptide 827. .832
FT /label= GXYGX'Y' _motif
FT 833. .838
FT /label= GXYGX'Y' _motif
FT 839. .844
FT /label= GXYGX'Y' _motif
FT 845. .850
FT /label= GXYGX'Y' _motif
FT 863. .868
FT /label= GXYGX'Y' _motif
FT 869. .874
FT /label= GXYGX'Y' _motif
FT 875. .880
FT /label= GXYGX'Y' _motif
FT 891. .896
FT /label= GXYGX'Y' _motif
FT 897. .902
FT /label= GXYGX'Y' _motif
FT 903. .908
FT /label= GXYGX'Y' _motif
FT 911. .916
FT /label= GXYGX'Y' _motif
FT 917. .922
FT /label= GXYGX'Y' _motif
FT 928. .933
FT /label= GXYGX'Y' _motif
FT 934. .939
FT /label= GXYGX'Y' _motif
FT 956. .961
FT /label= GXYGX'Y' _motif
FT 962. .967
FT /label= GXYGX'Y' _motif
FT 968. .973
FT /label= GXYGX'Y' _motif
FT 1126. .1131
FT /label= GXYGX'Y' _motif
FT 1145. .1150
FT /label= GXYGX'Y' _motif
FT 1193. .1198
FT /label= GXYGX'Y' _motif
XX
PN US5643783-A.
XX
PD 01-JUL-1997.
XX
PF 01-DEC-1993; 93US-00159784.
XX
PR 01-DEC-1993; 93US-00159784.
XX
PA (HARD) HARVARD COLLEGE.
XX
PI Olsen BR, Oh SP;
XX
DR WPI; 1997-350247/32.
DR N-PSDB; AAT84485.
XX
PT Nucleic acid encoding human alpha-1 collagen - for production of
PT recombinant alpha-1 collagen, for use in the treatment of cartilage
PT degeneration.
XX
PS Disclosure; Fig 2; 35pp; English.

Query Match 99.7%; Score 965; DB 2; Length 1288;
Best Local Similarity 99.5%; Pred. No. 5.4e-106;
Matches 183; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQARAVGLSGTFRFLSSRLQDLYSI 60
Db 1105 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQARAVGLSGTFRFLSSRLQDLYSI 1164
QY 61 VRRADRGSPVIVNLKDEVLSFSGSQGQVQPGARIFSFDRDVLHHPAWPQKSVW 120
Db 1165 VRRADRGSPVIVNLKDEVLSFSGSQGQVQPGARIFSFDRDVLHHPAWPQKSVW 1224

QY 121 HGSDPGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 1225 HGSDPGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 1284

QY 181 SFSK 184
Db 1285 SFSK 1288

RESULT 13
AAY18409
ID AAY18409 standard; protein; 184 AA.
XX AAY18409;
AC AAY18409;
XX
DT 24-AUG-1999 (first entry)
XX
DE Endostatin protein sequence.
XX
KW EM1; anti-angiogenic peptide; endostatin; angiogenesis-dependent cancer;
KW benign tumour; rheumatoid arthritis; psoriasis; ocular angiogenesis;
KW Osler-Webber Syndrome; myocardial angiogenesis; angiofibroma; cancer;
KW plaque neovascularisation; telangiectasia; atherosclerosis; scleroderma;
KW dialysis graft vascular access stenosis; renal cancer; therapy.
XX
OS Mus sp.
XX
XX WO9929855-A1.
PN
XX
PD 17-JUN-1999.
XX
PF 08-DEC-1998; 98WO-US026057.
XX
PR 08-DEC-1997; 97US-0067888P.
PR 22-APR-1998; 98US-0082663P.
PR 16-NOV-1998; 98US-0108536P.
XX
FA (BETH-) BETH ISRAEL DEACONESS MEDICAL CENT.
XX
XX Sukhatme VP;
PI
XX
DR WPI; 1999-385604/32.
DR N-PSDB; AAX79949.
XX
PT Mutant endostatin having anti-angiogenic activity.
XX
PS Claim 31; Fig 2; 105pp; English.
XX

This sequence is the mouse endostatin. The invention relates to a the mutant endostatin (EM), which has anti-angiogenic activity, and is designated EM1. Compositions comprising EM1 or fusion proteins comprising EM1, are useful for treating diseases characterised by angiogenic activity, such as angiogenesis-dependent cancers, benign tumours, rheumatoid arthritis, psoriasis, ocular angiogenesis, Osler-Webber Syndrome, myocardial angiogenesis, plaque neovascularisation, telangiectasia, haemophilic joints, angiofibroma, wound granulation, intestinal adhesions, atherosclerosis, scleroderma, hypertrophic scars, cat scratch disease, Helicobacter pylori ulcers, dialysis graft vascular access stenosis, contraception and obesity. In particular, the diseases treatable by EM1 comprise cancer, especially renal cancer. The methods provide a means for introducing EM1 into mammalian cells via gene therapy, for production of EM1 via recombinant means, as well as recombinant production of the EM1 protein. EM1 performs as well or better than whole endostatin. Use of EM1 is advantageous for treatment of angiogenic diseases in that increasingly smaller peptides are more potent on a weight basis, and may be able to better penetrate tissues

XX
SQ Sequence 184 AA;

Query Match 99.5%; Score 963; DB 2; Length 184;
Best Local Similarity 99.5%; Pred. NO. 5.3e-107;
Matches 183; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQOARAVGLSGTFRFLSSRLQDLYSI 60
Db 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFQOARAVGLSGTFRFLSSRLQDLYSI 60

QY 61 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFQGRDVLRHHPAWPKSVW 120
Db 61 VRRADRGSPVIVNLKDEVLSPSWDSLFSGSQQLQPGARIFSFQGRDVLRHHPAWPKSVW 120

QY 121 HGSDPGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180
Db 121 HGSDPGRRRLMESYCETWRTTGTATGQASSLLSGRLLLEQKAASCHNSYIVLCIENSFMT 180

QY 181 SFSK 184
Db 181 SFSK 184

RESULT 14
AAW92297
ID AAW92297 standard; peptide; 1288 AA.
XX AAW92297;
AC AAW92297;
XX
DT 28-APR-1999 (first entry)
XX
DE Mouse alpha-1 (XVIII) collagen chain common sequence MO18 (common) 28.
XX
KW Human; type XVIII collagen; liver disease; cirrhosis; detection;
KW hepatocellular carcinoma; diagnosis.
XX
OS Mus sp.
XX
XX WO9856399-A1.
XX
PD 17-DEC-1998.
XX
PF 12-JUN-1998; 98WO-US012327.
XX
PR 12-JUN-1997; 97US-0049369P.
XX
XX (FIBR-) FIBROGEN INC.
PA (FIFI-) ACAD FINLAND.
PA (INRM) INST NAT SANTE & RECH MEDICALE.
XX
PI Pihlajaniemi T, Rehn M, Clement B;
XX
DR WPI; 1999-070292/06.
XX

Diagnosis and monitoring of liver disease by measuring collagen type XVIII levels - with elevated levels indicative of disease, especially cirrhosis or hepatocellular carcinoma.

Example 6; Fig 8; 56pp; English.

A method has been developed for the detecting liver disease. The method comprises: (a) reacting a patient sample with antibodies (Ab) specific for collagen type XVIII (Col18); (b) measuring the amount of Ab-antigen complex (C) formed as indicator of the amount of Col18 present; (c) similar analysis of a non-diseased control; and (d) comparing the amounts of Col18 in the two samples to detect presence or progression of disease. Elevated levels of Col18 are: (i) indicative of disease, specifically cirrhosis; and (ii) predictive of the prognosis of disease, specifically hepatocellular carcinoma (there is a relationship between Col18 mRNA levels and tumour size and necrosis, and survival times are significantly higher in patients with higher Col18 levels). The method provides non-invasive, early and accurate diagnosis of liver disease. The present sequence represents the sequence common to mouse alpha-1 (XVIII) collagen chain from the present invention

XX
SQ Sequence 1288 AA;

Query Match 99.2%; Score 960; DB 2; Length 1288;

Best Local Similarity 99.5%; Pred. No. 2.2e-105;
Matches 182; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFOQARAVGLSGTFRFLSSRLQDLYSI 60
Db 1106 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFOQARAVGLSGTFRFLSSRLQDLYSI 1165
QY 61 VRRADRGSPVIVNLKDEVLSFSGQQLPGARIFSFDRDVLRRHPAPQKSVW 120
Db 1166 VRRADRGSPVIVNLKDEVLSFSGQQLPGARIFSFDRDVLRRHPAPQKSVW 1225
QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEOKAASCHNSYIVLCIENSEFMT 180
Db 1226 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEOKAASCHNSYIVLCIENSEFMT 1285
QY 181 SFS 183
Db 1286 SFS 1288

RESULT 15

AAAY25114
ID AAAY25114 standard; protein; 684 AA.

XX AC AAAY25114;

XX DT 25-AUG-1999 (first entry)

XX DE Mouse alpha1 (XVIII) collagen protein.

XX KW Alpha(XVIII) collagen; mimetic; endostatin; atomic coordinate; library;
KW anti-angiogenic; heparin binding domain; receptor binding domain; mimic;
KW alpha-helix A domain; carbohydrate recognition domain; CRD domain;
KW treatment; angiogenesis; tumour; murine.

XX OS Mus sp.

XX PN WO9931616-A1.

XX PD 24-JUN-1999.

XX PF 16-DEC-1998; 98WO-US026783.

XX PR 16-DEC-1997; 97US-0069727P.

XX PA (HARD) HARVARD COLLEGE.

XX PI Olsen BR, Hohenester E, Timpl R, Sasaki T;

XX DR WPI; 1999-395243/33.

XX PT Identifying mimetics of mammalian endostatin.

XX PS Disclosure; Fig 5A-C; 75pp; English.

XX CC This invention describes a novel method for identifying mimetics of
XX CC mammalian endostatin. The method comprises identifying a compound having
XX CC atomic coordinates with non-trivial similarity to selected coordinates of
XX CC atoms of a mammalian endostatin involves (a) providing a library of
XX CC atomic coordinates of compounds in a library of candidate compounds, (b)
XX CC comparing the library of atomic coordinates to the selected coordinates
XX CC of a mammalian endostatin and (c) selecting from the library at least one
XX CC candidate compound on the basis of selection criteria which include
XX CC similarities between the atomic coordinates of the selected candidate
XX CC compound and the atomic coordinates of the mammalian endostatin. The
XX CC invention also describes the use of an anti-angiogenic fragment of
XX CC endostatin comprising a domain selected from a heparin binding domain, a
XX CC receptor binding domain, and exposed on alpha-helix A domain, and a
XX CC carbohydrate recognition domain (CRD) domain. The methods can be used for
XX CC designing and selecting endostatin mimics. The compounds identified can
XX CC be used for treating undesired angiogenesis, e.g. tumours. This sequence
XX CC represents mouse alpha1(XVIII) collagen which is used in the description
XX CC of the method

XX SQ Sequence 684 AA;

Query Match 98.1%; Score 950; DB 2; Length 684;
Best Local Similarity 99.4%; Pred. No. 1.4e-104;
Matches 180; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFOQARAVGLSGTFRFLSSRLQDLYSI 60
Db 502 HTHQDFQPVHLVALNTPLSGGMRGIRGADFQCFOQARAVGLSGTFRFLSSRLQDLYSI 561
QY 61 VRRADRGSPVIVNLKDEVLSFSGQQLPGARIFSFDRDVLRRHPAPQKSVW 120
Db 562 VRRADRGSPVIVNLKDEVLSFSGQQLPGARIFSFDRDVLRRHPAPQKSVW 621
QY 121 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEOKAASCHNSYIVLCIENSEFMT 180
Db 622 HGSDPSGRRRLMESYCETWRTTGTGATGQASSLLSGRLLEOKAASCHNSYIVLCIENSEFMT 681
QY 181 S 181
Db 682 S 682

Search completed: August 31, 2004, 19:55:23
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